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None

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B1H
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(54) Chromatographic separation device

(57) A chromatographic separation device comprises a body 2 of a semiconductor material which body has a longitudinal channel 1 formed in a surface thereof, the channel 1 being capable of containing a predetermined volume of a liquid or solid material for a chromatography test or separation procedure, the channel carrying at least one electrode 6 positioned intermediate the channel ends. The semiconductor body may additionally support an electronic or optical sensor 8 arranged in line with said channel 1 to provide an integrated detection system.

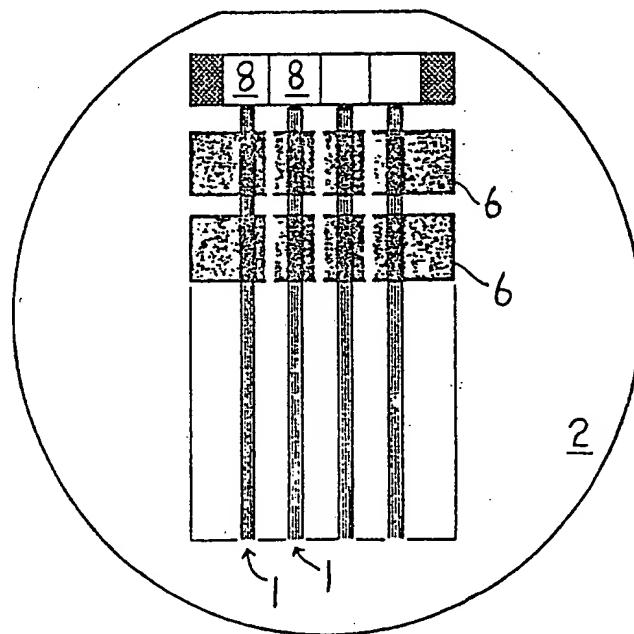


Fig. II

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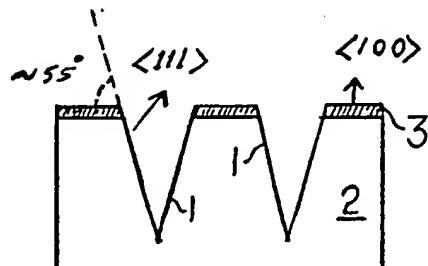


Fig.1

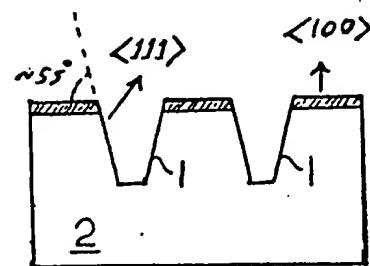


Fig.2

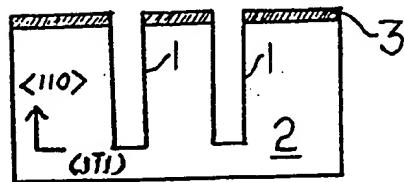


Fig.3

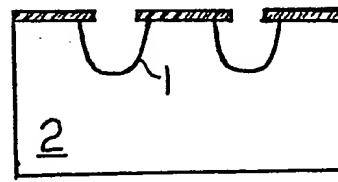


Fig.4



Fig.5

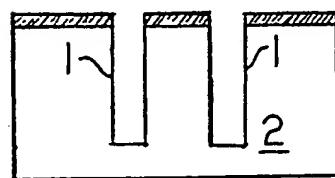


Fig.6

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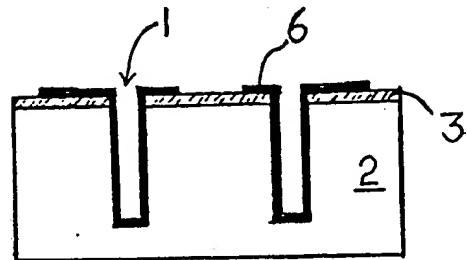


Fig. 7

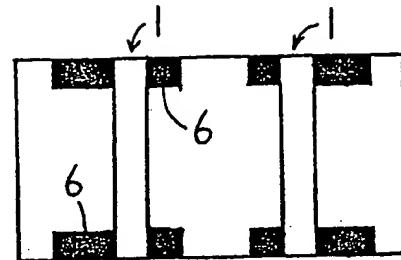


Fig. 9

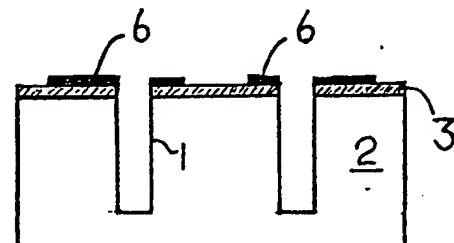


Fig. 8

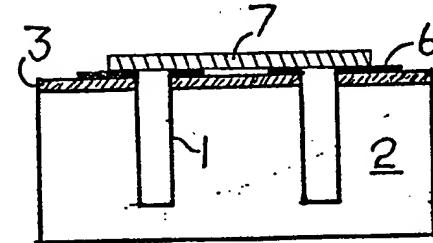


Fig. 10

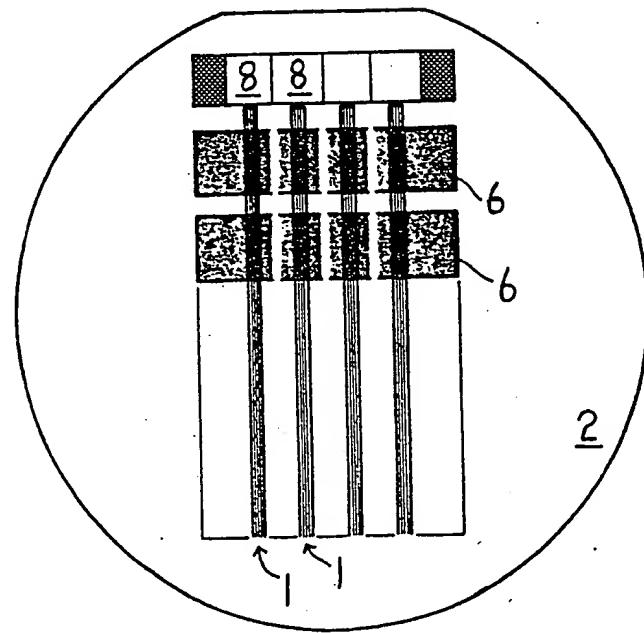


Fig. 11

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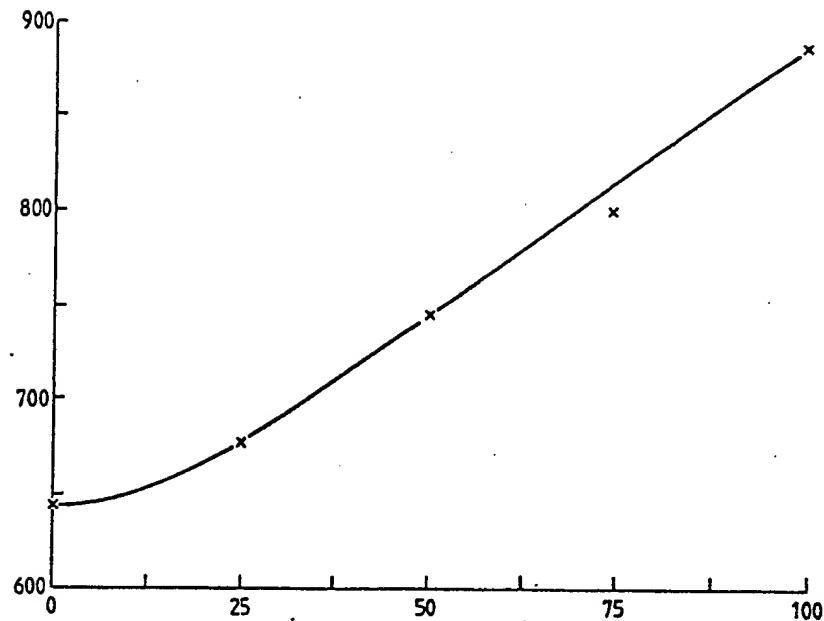


Fig.12

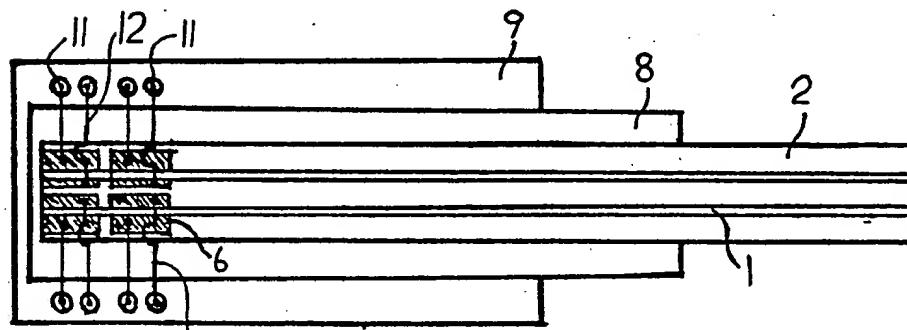


Fig.13

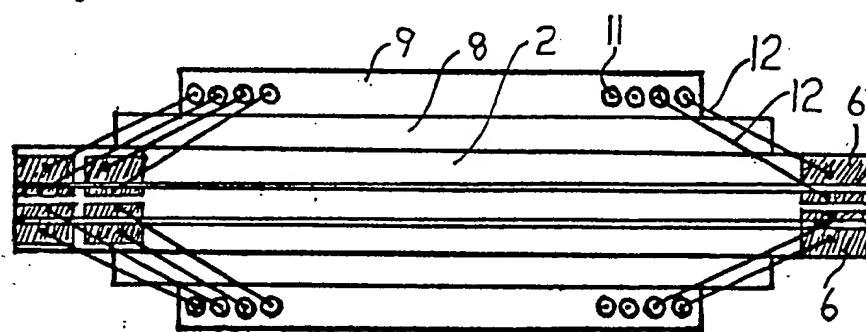


Fig.14

SPECIFICATION

Chromatographic separation device

5 This invention relates to a chromatographic separation device. It relates particularly to the provision of such a device that can be used for a variety of applications such as electrolysis, chromatography, electrophoresis and the study of electrokinetic phenomena.

10 The development of separation technology has brought with it the need to be able to work accurately with very small test samples and possibly to provide a separation cell that can be integrated with a detection system. The signals from an integrated sensor will be less likely to be influenced by noise or leakage as compared with those of a discrete sensor. The sensing element would thus make use of a non-specific technique such as conductivity, optical (absorbance, refractive index) or electrochemical properties or, more likely, a suitable combination of these.

15 The present invention was devised to provide a separation device that was capable of being manufactured in a miniature form to assist the microseparation and detection of biochemical and other chemical species.

20 According to the invention, there is provided a chromatographic separation device comprising a body of a semiconductor material which body has a longitudinal channel formed in a surface thereof, the channel being capable of containing a predetermined volume of a liquid or solid material for a chromatographic test or separation procedure, the 25 channel carrying at least one electrode positioned intermediate the channel ends.

30 Preferably, an open side of the said channel is closed by a cover plate.

35 The channel may be formed by an integrated circuit technique such as photolithography or micromachining. Alternatively, the channel may be formed by a micromechanical machining technique such as electromechanical sawing.

40 The body of semiconductor material may be a silicon wafer. The separation device may further carry an electronic or optical sensor element which is located in line with the channel. The body of semiconductor material may be provided with two or more of the longitudinal channels, the channels 45 being located in a mutually parallel arrangement.

45 By way of example, some particular embodiments of the invention will now be described with reference to the accompanying drawings, in which:-

50 *Figure 1* is a cross-sectional view greatly enlarged 55 of an anisotropically etched microchannel structure in a body of silicon wafer material;

55 *Figure 2* is a similar view showing a different channel shape;

60 *Figure 3* is a similar view showing the result of etching a body of silicon having a different crystal orientation;

65 *Figure 4* is a similar view showing the result of an isotropic etching process;

70 *Figure 5* is a plan view showing the formation of a 65 channel having a serpentine shape by the etching

process;

75 *Figure 6* is similar to *Figure 4* and shows the result of an electromechanical sawing process;

80 *Figure 7* is a cross-sectional view showing a metal contact pattern deposited on a channel;

85 *Figure 8* is a similar view showing the metal contact pattern having been etched or sawed to partially remove the metal deposit;

90 *Figure 9* is a plan view of the electrode structure depicted in *Figure 8*

95 *Figure 10* is a cross-sectional view of the electrode structure on a channel with a cover plate placed over the channel;

100 *Figure 11* is a plan view of a silicon wafer provided with four chromatographic separation devices of the invention and an array of electronic or optical sensors;

105 *Figure 12* is a graph showing the results of some conductance measurements made using the separation devices;

110 *Figure 13* is a plan view showing a prototype construction of a single ended chromatographic separation device, and,

115 *Figure 14* is a similar view of a double ended separation device.

120 The construction of the chromatographic separation device of the invention begins with the preparation of a slice of semiconductor material and in the embodiment about to be described this was a silicon wafer. For convenience in use of the separation device some of the test slices were formed with two channels located in a mutually parallel arrangement.

125 The formation of very narrow channels in silicon wafer material can be achieved by using the well-developed integrated circuit technology of photolithography, micromachining or micromechanical machining. Micromachining utilises the controlled etching characteristics of silicon, and involves anisotropic and isotropic wet and/or dry etching. Aniso-

130 tropic etchants, which are also known as orientation-dependent or crystallographic etchants, etch silicon at different rates in different directions in the crystal lattice; they can form a variety of well-defined shapes with sharp edges and corners. Typical examples include hot alkaline solutions such as aqueous potassium hydroxide (KOH) or sodium hydroxide and a mixture of ethylenediamine, pyrocatechol and water known as EDP. Dry etching techniques such as reactive ion etching and argon ion beam mil-

135 ling can also be employed to perform anisotropic etching. Isotropic etchants, on the other hand, etch the silicon crystal at the same rate in all directions and generally produce rounded shapes. Typical examples include mixtures of hydrofluoric, nitric and acetic acids known as HNA.

140 The fabrication of the channel structures by micromachining involved the following main steps: formation of a layer of silicon dioxide on the silicon wafer body by a standard thermal oxidation process; definition of patterns on the oxidised surface by using standard photoresist and photolithography processes; removal of oxide by wet or dry etching where channels were to be formed thus exposing the silicon surface; removal of photoresist and etching the silicon body in places where it was unprotected by

the oxide mask. In structures requiring long etching times, silicon nitride was generally employed as a masking material in place of silicon oxide (SiO_2). In certain cases, gold and chromium metals might 5 alternatively be employed for this purpose.

Anisotropic wet etching was performed mainly on silicon wafers with two alternative types of crystal surface orientations viz. $\langle 100 \rangle$ and $\langle 110 \rangle$. Etching along $\langle 110 \rangle$ planes was quite rapid compared with 10 $\langle 100 \rangle$ planes. The attack along a $\langle 111 \rangle$ plane was extremely slow, if it occurred at all, by the action of anisotropic etchants. A variety of channels was formed by controlling orientation, shape and size of the oxide openings on the surface of these wafers, and 15 etching silicon with standard anisotropic etchants mentioned earlier. When etching with these etchants, proper mask alignment with specific crystallographic axes of the wafer was considered of utmost importance if the required structures were to 20 be precise. Some typical cross-sectional profiles of channels etched in this way in silicon for utilising in chromatographic and electrophoretic devices are described below.

As shown in Figure 1, V-shaped channels 1 were 25 formed in a silicon wafer body 2 by an alkaline etchant (KOH) acting through rectangular openings in an oxide mask 3 oriented along the $\langle 110 \rangle$ direction of $\langle 100 \rangle$ wafer with $\langle 111 \rangle$ side walls. Etching was stopped in the early stages to produce the structure 30 shown in Figure 2. In the formation of narrow channels, the depth needed to be closely controlled by the width of the opening in the oxide etch mask. For producing deeper structures (50 microns), silicon nitride was employed as the masking material. Electro- 35 chemical etching was also employed to reduce the problem of undercutting the mask. In a $\langle 110 \rangle$ oriented wafer, two sets of $\langle 111 \rangle$ planes are aligned perpendicular to the $\langle 110 \rangle$ surface plane although not to each other. Long, deep and closely spaced channels, 40 with vertical wall $\langle 111 \rangle$ side terminations (Figure 3), were etched by potassium hydroxide reagent in a $\langle 110 \rangle$ silicon wafer; the etching ratio in the $\langle 110 \rangle$ to $\langle 111 \rangle$ direction was very high (~400 to 1).

An isotropic etchant (HNA), with continuous agitation, was employed to produce the channel structures shown in Figure 4. The HNA etch can be employed to produce a variety of channel structures such as serpentine, spiral, etc. the alignments of which are independent of crystallographic orientations of the silicon wafer material. A plan view of a typical serpentine etched structure is shown in Figure 5.

Although rarely used for etching silicon, the technique of micromechanical machining has been 55 found to be suitable as an alternative process for forming the channels. It mainly involves electro-mechanical sawing with very fine diamond blades under controlled conditions to produce nearly rectangular, closely spaced channels in the silicon. The 60 shapes formed are independent of crystallographic orientation of the silicon wafer. A typical sawed cross section (75 microns \times 200 microns) is shown in Figure 6.

The metal contact electrodes required in the fabrication of electrophoretic devices and for the detection

sensing of species by conductivity and other electrochemical measurements were generally deposited by a standard multimetal sputter deposition process. Four alternative metallisation schemes

70 viz. titanium/gold, titanium/platinum/gold, chromium/gold and chromium/platinum/gold were employed for this purpose. The thickness of each of the metals titanium, platinum and chromium was usually around 100nm whereas that of gold varied 75 between 1 micron to 3 microns depending upon particular design and application. Metal contact patterns were produced either by using standard photoresist, photolithography, metal etching and resist float-off technique or by depositing metals directly through 80 contact ceramic or metal masks. A variety of etched and sawed column structures were electrode; one typical cross section view of such a structure is given in Figure 7.

As depicted in Figure 7, the silicon wafer body 2 85 carries a silicon oxide mask 3 having a thickness of about 0.5 microns and above the channel 1 an electrode 6 area has been deposited. The electrode 6 area covers the two sides of the channel 1 and it also descends down the side walls of the channel and across 90 the channel bottom. The presence of electrode material in the channel is not required because it reduces the cross-sectional area of the channel. Accordingly, this excess electrode material was removed by etching or by the electromechanical sawing operation to 95 leave only the portions of electrode 6 which lie on top of the oxide mask 3.

The cross-sectional view of the wafer body 2 then 100 appears as in Figure 8, and a plan view is given in Figure 9. A view of the wafer body with the cover plate 7 in place is given in Figure 10.

Figure 11 gives a plan view of a practical construction in which a circular silicon wafer body 2 having a diameter of three inches forms a substrate for four channels 1 each having a width about 75 microns 105 and a depth of about 200 microns. Each channel 1 has two electrodes 6, the two electrodes being spaced longitudinally along the length of the channel. At the end of each channel 1 there is provided an electronic or optical sensor 8, the four sensors being 110 mounted as an array on the surface of the wafer body 2.

The resulting open channel structures formed by the abovementioned techniques in silicon wafer material were covered over with a Pyrex (Registered 115 Trade Mark) glass cover plate which was merely placed over the open side of the channels or in alternative embodiments was bonded into place anodically or with an adhesive. This produced an enclosed capillary channel structure the dimensions of which 120 could be varied from a fraction of a micron to many microns (about 300 microns). The minimum dimensions of the channels which can be employed in chromatographic devices will be limited by the 'molecular size' of the species under detection 125 whereas in the case of electrophoretic devices these will be mainly controlled by the thickness of the double layer of the species. The nonaqueous systems such as proteins and macromolecules are most likely to have larger values of the double layer thickness (a few orders of magnitude) compared with 130

aqueous systems (~5 A.U.).

Use of the chromatographic separation device of the invention is illustrated by the following Examples:

5

Example 1. Chromatography in microchannels and detection on a silicon wafer

In a sample of the device corresponding to that shown in Figure 11, the channels (width ~75 microns, depth ~200 microns) were filled with a 1.5% agarose solution, immobilised and held in place by covering with a glass cover plate. The operation of filling the channel was effected by conventional methods such as diffusion and the application of 15 high pressures. The wafer was placed in a beaker containing a small quantity of the enzyme β -galactosidase in sodium phosphate buffer solution. The device was then left standing overnight in a vertical position with the lower channel ends just 20 dipping in the solution. The channels were subsequently analysed for enzyme activity using 4-methylumbelliferyl- β -D-galactopyranoside, which is enzymatically cleaved to form a fluorescent product. Under UV irradiation the channels were seen to 25 fluoresce and it was concluded that the enzyme had migrated up the channels.

To detect the species near the 'exit' end of the channels by conductivity measurements, the above experiment was repeated using sodium phosphate 30 buffer solutions of various concentrations. Agarose was immobilised in the channels as before. The devices were allowed to stand upright in a small volume of sodium phosphate buffer solution, and the conductance between the two metal pad electrodes was 35 then measured. The results are shown in the graph of Figure 12, where the vertical axis measures conductance (in microseconds) and the horizontal axis measures concentration (in millimoles) of the relevant sodium phosphate buffer solution. It can be 40 seen that the steady state conductance is linearly related to the ionic strength of the buffer solution.

These two experiments demonstrate that it is possible for biochemical or other chemical species to migrate along these microscopic channels and to be 45 detected fluoroscopically or for the concentration of the migrating chemical to be determined quantitatively by the conductance measurements when using an integrated sensor.

50 *Example 2. A chromatographic separation and sensing device*

To construct a practical chromatographic device as shown in Figure 13, a portion of the silicon wafer body 2 measuring 7mm x 65mm and being provided 55 with rectangular channels 1 (dimensions: 100 microns x 200 microns) was mounted using an epoxy resin adhesive onto alumina strips 8. The alumina strips 8 measured 13mm x 50mm. In turn, the strips 8 were epoxy resin bonded to a gold plated electrodes 60 mounting package 9 having insulated pins 11 for making electrical connections. The chromatographic devices were then bonded to the pins 11 by means of gold wires 12 attached to the electrode 6 areas.

In tests carried out on the separation and sensing 65 properties of this device, the channels were filled

with a variety of chromatographic media for the resolution of gases and volatiles by gas chromatography and a variety of other materials by liquid chromatography. Thus, chromatographic media

70 comprising styrene/divinylbenzene ion exchange resins and acrylic carboxylic, tertiary amino and chelating resins were formed *in situ* by polymerisation onto an allyldimethyl chlorosilane-activated channel. These media were exploited for the resolution of a variety of charged metallic, organic and proteinaceous species. Similarly, silica or alumina adsorbents comprising oxidised silicon or aluminium channels were used for the micro-chromatography of complex lipids, fatty acids, 75 steroids, alkaloids, phenols, hydrocarbons, dicarboxylic acids, amino acids, esters, peroxides, aldehydes, alcohols and nucleic acids.

Proteins and enzymes were resolved by gel filtration media comprising sephadexes, agaroses, acrylics or porous silicas, by ion exchange on carbohydrate-based or acrylic exchanges, hydrophobic adsorbents such as phenyl or alkyl-agaroses. Selective adsorption of individual proteins or groups of proteins was achieved by micro-affinity chromatography on immobilised dyes, lectins, chelating ligands, protein A, boronate, heparin, nucleotides, immunoligands etc. Similarly, charge transfer ligands were used to resolve nucleotides, immunoligands etc. Similarly, charge transfer ligands 95 were used to resolve nucleotides, oligonucleotides and nucleic acids, and appropriate chiral phases for the resolution of isomeric and chiral compounds.

Example 3. An electrophoretic separation and sensing device

In a similar construction to that used for the chromatographic device of Example 2, an electrophoretic device as depicted in Figure 14 was built. For the electrophoretic device, the electrode 6 areas 105 needed to be provided at both ends of the channels 1; consequently the pins 11 and the gold wires 12 were similarly required to be located at both the ends of the channels 1.

In tests carried out on the separation and sensing 110 properties of the device, it was found that the positioning of the electrodes at the distal ends of the channel 1 filled with an appropriate electrophoretic medium permitted miniaturised electrophoresis to be performed within minutes. For example, electrophoresis

115 in agarose gel resolved proteins according to charge, whilst polyacrylamide gel electrophoresis was found to impose a sieving effect additional to the charge separation. Such a miniaturised device containing agarose or acrylamide gels could be used to 120 resolve proteins, enzymes and isoenzymes in normal and abnormal serum samples.

Similarly, agarose and polyacrylamide gel electrophoresis may be used for the resolution of nucleic acids and oligonucleotides and the device finds particular application in DNA restriction fragment analysis, DNA sequencing and probe analysis. These supports are also applicable to a variety of immunological, electrofocussing and affinity techniques.

130 The chromatographic separation device of the in-

vention is proposed to be employed to study the electrokinetic or zeta potential, which is involved in electro-osmosis, electrophoresis and allied phenomena. The zeta potential is the potential between the fixed and freely mobile part of the double layer and for a certain class of electrolytes it has been generally reported to control the electrophoretic mobility. A knowledge of these parameters such as the zeta potential of certain ionic species under examination may be of help to predict the response speed of these ions for detection.

The foregoing description of embodiments of the invention has been given by way of example only and a number of modifications may be made without departing from the scope of the invention as defined in the appended claims. For instance, in some applications it may be possible to use the separation device without need for a cover plate over the open side of the channel.

20 CLAIMS

1. A chromatographic separation device comprising a body of a semiconductor material which 25 body has a longitudinal channel formed in a surface thereof, the channel being capable of containing a predetermined volume of a liquid or solid material for a chromatographic test or separation procedure, the channel carrying at least one electrode positioned intermediate the channel ends.
2. A separation device as claimed in claim 1, in which an open side of the said channel is closed by a cover plate.
3. A separation device as claimed in claim 1 or 2, 35 in which the said channel is formed by an integrated circuit technique such as photolithography or micro-machining.
4. A separation device as claimed in claim 1 or 2, in which the said channel is formed by a micro-40 mechanical machining technique such as electro-mechanical sawing.
5. A separation device as claimed in any one of claims 1 to 4, in which the said body of semiconductor material is a silicon wafer.
- 45 6. A separation device as claimed in any one of claims 1 to 5, in which the said body further carries an electronic or optical sensor element located in line with said channel.
7. A separation device as claimed in any one of 50 claims 1 to 6, in which the said body is provided with two or more of said longitudinal channels, the channels being located in a mutually parallel arrangement.
8. A chromatographic separation device substantially as hereinbefore described with reference 55 to any one of the accompanying drawings.